

Supplier's self-disclosure

Quality - Environment – Health and safety



Supplier:	Address:		
Email:	Phone:	Fax:	
Product range:			
References:			

Responsibilities	Name	Capacity	Extension
Commercial management:			
Technical management:			
Quality:			
Environment:			
Health and safety:			
Sales:			
Purchase:			
Production:			
Total number of employees:		Thereof in the production:	
Turnover for the last 3 years:	20__:	20__:	20__:
Do you have a product-liability insurance?		yes	no

Do you have a certified:

Quality Management System (QMS)?		Norm:	Certifier:	Reg.-No.:	Valid until:
	yes*				
	no				
Environmental Management System (UMS)?		Norm:	Certifier:	Reg.-No.:	Valid until:
	yes*				
	no				
Health and safety Management (AMS)?		Norm:	Certifier:	Reg.-No.:	Valid until:
	yes*				
	no				

* Please send us a copy of the certificate(s).

If all questions concerning the relevant management systems were answered „yes“ the processing of the following questionnaire is not required.

The supplier's self-disclosure has been edited by:

Mrs / Mr:

Capacity:

Place and date

Signature

Questionnaire

1. Organization					
1.1	Does your company have a documented QM system	yes	no	According to which standard?	Since when?
	QM - System?				
	UM - System?				
	AS - System?				
1.2	Is certification planned?	yes	no	expected date?	
	QM - System:				
	UM - System:				
	AS - System:				
1.3	Is a division „quality assurance“ provided?				
1.4	Is this division directly responsible to the management?				

2. Contract review			yes	no
2.1	Are procedures and responsibilities laid down for order processing?			
2.2	Are these clearly regulated by method or process description?			

3. Development			yes	no
3.1	Do you practice your own product development?			
3.2	Is the development process documented?			
3.3	When developing do you also consider safety and environmental aspects?			

4. Documentation			yes	no
4.1	Is there a distribution and revision system for drawings, standards, specifications, safety and environmental regulations etc.?			
4.2	Are procedures and responsibilities clearly regulated in writing?			

5. Procurement			yes	no
5.1	Are procedures and responsibilities for procurement set out in writing?			
5.2	Are also environmental and security-relevant aspects taken into account in the procurement?			
5.3	Is a procedure for the selection, release and evaluation of suppliers set?			

6. Supplied products by contracting authorities			yes	no
6.1	Are there regulations for dealing with customer-supplied products / services?			
6.2	Is the liability for supplied products, tools, forms etc. regulated?			

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7. Labelling and traceability		yes	no
7.1	Are there any regulations ensuring the identifiability of products and the assignment to the appropriate contract documents?		

8. Production		yes	no
8.1	Are procedures and responsibilities for production specified in writing?		
8.2	Are there process descriptions or procedures that govern the flow of production, installation, maintenance etc.?		
8.3	Is it ensured that the implementation of the operations and testing activities will be documented?		
8.4	Is a preventive maintenance of production equipment carried out?		

9. Quality inspection		yes	no
9.1	Are procedures and responsibilities for incoming goods, intermediate and final testing set out in writing?		
9.2	Are there process descriptions or procedures available?		
9.3	Are the results of a test recorded?		

10. Inspection of measuring and testing equipment		yes	no
10.1	Is there a procedure set for the inspection of measuring and testing equipment?		
10.2	Are all measuring and testing equipment systematically identified?		

11. Inspection status		yes	no
11.1	Is the inspection status of raw materials, semi-finished and finished products visible at all times?		

12. Control of non-conforming products		yes	no
12.1	Are non-compliant products identified?		
12.2	Is described how to record, store and report defective parts?		
12.3	Is the customer always informed about product and schedule variances?		

13. Corrective and preventive measures		yes	no
13.1	Are methods described to guarantee the elimination of causes of non-conformance in case of process or product variances?		
13.2	Are non-conformance and variances recorded in writing and analyzed?		
13.3	Are near accidents and accidents recorded and analyzed and corrective actions implemented?		
13.3	Are environmental events recorded and analyzed and corrective actions implemented?		

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14. Handling, storage, packing, labelling and shipping		yes	no
14.1	Is the handling, packing, labelling, storage, shipping and the load securing set out in writing?	<input type="checkbox"/>	<input type="checkbox"/>
14.2	Are there any procedures and work instructions available?	<input type="checkbox"/>	<input type="checkbox"/>

15. Quality records		yes	no
15.1	Has it been determined what quality records should be maintained and what retention periods apply?	<input type="checkbox"/>	<input type="checkbox"/>

16. Training		yes	no
16.1	Is the qualification for persons with quality-related activities specified in writing?	<input type="checkbox"/>	<input type="checkbox"/>
16.2	Are instructions on the subject of quality, environmental protection and occupational safety documented?	<input type="checkbox"/>	<input type="checkbox"/>

17. After-sales service		yes	no
17.1	Is the handling and processing of customer complaints regulated in writing?	<input type="checkbox"/>	<input type="checkbox"/>